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=> s Kontsekova E?/au
            48 KONTSEKOVA E2/AU
=> s 11 and tau
         87233 TAU
           168 TAUS
         87285 TAU
                 (TAU OR TAUS)
L2
            10 L1 AND TAU
=> s 12 and truncated
        41439 TRUNCATED
             4 L2 AND TRUNCATED
=> d ibib abs 1-4
   ANSWER 1 OF 4 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER:
                       2006:607535 CAPLUS
DOCUMENT NUMBER:
                         145:121973
TITLE:
                         Truncated tau from sporadic
                         Alzheimer's disease suffices to drive neurofibrillary
                         degeneration in vivo
AUTHOR(S):
                         Zilka, Norbert; Filipcik, Peter; Koson, Peter;
                         Fialova, Lubica; Skrabana, Rostislav; Zilkova, Monika;
                         Rolkova, Gabriela; Kontsekova, Eva; Novak,
                        Michal
CORPORATE SOURCE:
                        Axon Neuroscience GmbH, Vienna, 1030, Austria
SOURCE:
                        FEBS Letters (2006), 580(15), 3582-3588
                        CODEN: FEBLAL: ISSN: 0014-5793
PUBLISHER:
                        Elsevier B.V.
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
    Truncated tau protein is the characteristic feature of
AB
     human sporadic Alzheimer's disease. We have identified truncated
     tau proteins conformationally different from normal healthy
     tau. Subpopulations of these structurally different tau
     species promoted abnormal microtubule assembly in vitro suggesting toxic
     gain of function. To validate pathol. activity in vivo we expressed
     active form of human truncated tau protein as
     transgene, in the rat brain. Its neuronal expression led to the
     development of the neurofibrillary degeneration of Alzheimer's type.
     Furthermore, biochem, anal, of neurofibrillary changes revealed that
     massive sarcosyl insol. tau complexes consisted of human
     Alzheimer's tau and endogenous rat tau in ratio 1:1
     including characteristic Alzheimer's disease (AD)-specific proteins (A68).
     This work represents first insight into the possible causative role of
     truncated tau in AD neurofibrillary degeneration in
     wive
REFERENCE COUNT:
                               THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS
                         30
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
    ANSWER 2 OF 4 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER:
                         2004:60551 CAPLUS
DOCUMENT NUMBER:
                         140:124832
                        Truncated tau proteins
TITLE:
INVENTOR(S):
                        Kontsekova, Eva
PATENT ASSIGNEE(S):
                       Axon Neuroscience Forschungs- und Entwicklungs GmbH,
                        Austria
SOURCE .
                        PCT Int. Appl., 97 pp.
                        CODEN: PIXXD2
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DOCUMENT TYPE: Patent LANGUAGE: English FAMILY ACC. NUM. COUNT: 2 PATENT INFORMATION:

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						A2				WO 2003-EP7389										
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												, KG,								
												, MW,								
												, SG, , YU,				TJ,	TM,	TN,		
		PW-										, IU,				ΔM	17	BY		
		1411.										, CH,								
												, NL,								
												, GW,								
	AU 2003253044			A1 20040202				AU 2003-253044 EP 2003-763763					20030709							
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	EP	1521																		
		R:										, IT,						PT,		
												, TR,								
		1668				A		2005	0914		CN :	2003-	8166	47		2	.0030	709		
	JΡ	2006 3917 4063	5152	70		T		2006	0525		JP :	2004-	5205	41		2	.0030	709		
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	ΑT	4063	83			T		2008	0915		AT :	2003-	7637	63		2	.0030	709		
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	EP	1995	255			A1		2008	1126		EP :	2008-	1470	6		2	0030	709		
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			IT,	LI,	LU,	MC,	NL,	PT,	RO,	SE,	SI	, SK,	TR,	LT,	LV					
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	US	2006	0167	227		A1		2006	0727		US :	2005-	5211	40		2	0051	031		
PRIO	IORITY APPLN. INFO.:							US 2005-521140 AT 2002-1053						A 2	0020	712				
											EP :	2003-	7637	63		A3 2	0030	709		
											WO :	2003-	EP73	89		W 2	0030	709		
AB	Des	crib	ed a	re n	ovel	N-	and	C-te	rmin	ally	do	uble	trun	cate	d					

tau mols., (type IA, IB, IIA and IIB tau mols.) as well as methods for providing these mols., both from recombinant and biol. sources. Moreover, screening methods using these mols. in connection with Alzheimer's diagnosis and therapy are provided.

REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD, ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2009 ACS on STN 2003:64022 CAPLUS

ACCESSION NUMBER:

DOCUMENT NUMBER: 138:367296

TITLE: DC11: a novel monoclonal antibody revealing

Alzheimer's disease-specific tau epitope Vechterova, Lubica; Kontsekova, Eva; Zilka, AUTHOR(S):

Norbert; Ferencik, Miroslav; Ravid, Rivka; Novak, Michal

Axon Neuroscience, Vienna, A-1030, Austria CORPORATE SOURCE:

NeuroReport (2003), 14(1), 87-91 SOURCE: CODEN: NERPEZ; ISSN: 0959-4965

PUBLISHER: Lippincott Williams & Wilkins DOCUMENT TYPE: Journal LANGUAGE: English

AB Using tau protein exts. from Alzheimer's disease (AD) brain

tissue, we generated a monoclonal antibody (mAb DC11) which decorated

neurofibrillary pathol. in brain derived from AD patients on immunohistochem., and which lacked reactivity with healthy brain tissue. The same pattern of DCII specificity was observed on Western blot. The main constituent of structures decorated by DCII is microtubule-associated protein tau. In Western blot, DCII recognized neither native healthy tau nor its full length recombinant counterpart. However, the mAb showed strong immunoreactivity with truncated tau (residues tau.151-421), thus indicating the requirement for a

conformational epitope. Importantly, the DC11 epitope was phosphorylation independent. The immunochem. parameters of mAb show that DC11 could represent a novel structural probe with the specificity for conformation of pathol. tau present in Ab brains.

REFERENCE COUNT: 20 THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1995:814757 CAPLUS
DOCUMENT NUMBER: 123:250505

ORIGINAL REFERENCE NO.: 123:44639a,44642a
TITLE: Quick purification of recombinant human

truncated tau proteins for

immunoanalysis

AUTHOR(S): Kontsekova, Eva; Cattaneo, Antonino; Novak,

Michal

CORPORATE SOURCE: Institute of Virology, Slovak Academy of Sciences, 842

46, Bratislava, Czech.

SOURCE: Journal of Immunological Methods (1995), 185(2), 245-8

CODEN: JIMMBG; ISSN: 0022-1759

PUBLISHER: Elsevier DOCUMENT TYPE: Journal LANGUAGE: English

AB A simple and rapid purification method is described which exploits the heat stability of human tau (.tau.) protein to prepare

truncated forms of this protein derived from bacteria. Bacterial

cells expressing .tau. fragments were pelleted, resuspended in phosphate buffered saline and boiled for 5 min. After centrifugation the

supernatant containing thermostable .tau. was filtered (0.45  $\mu m)$ 

and used for immunoanal. with monoclonal antibodies. The purified . tau. fragments exhibited identical antigenic properties as

fragments isolated by a conventional procedure, based on ion exchange

chromatog. on phosphocellulose. In contrast to the conventional approach, our method is less complicated, cheaper and significantly reduces the time recuired for isolation of the recombinant tau. fragments.

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E1 THROUGH E27 ASSIGNED

=> d all 13 2

L3 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2009 ACS on STN

AN 2004:60551 CAPLUS

DN 140:124832

ED Entered STN: 26 Jan 2004

TI Truncated tau proteins

IN Kontsekova, Eva

PA Axon Neuroscience Forschungs- und Entwicklungs GmbH, Austria

SO PCT Int. Appl., 97 pp. CODEN: PIXXD2

DT Patent

LA English

CC 9-2 (Biochemical Methods)

Section cross-reference(s): 3, 6, 13, 14, 15

FAN.CNT 2

FAN.CNT 2	KTND	Dame.	ADDITOARTON NO	D									
PATENT NO.	KIND	DAIE	APPLICATION NO.										
PI WO 2004007547 WO 2004007547	A2 A3	20040122 20040722	WO 2003-EP7389	20030709									
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EP 1521774 R: AT. BE.	B1	20080827	, GR, IT, LI, LU,	NI CE MO DE									
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CN 1668641	A	20050914	CN 2003-816647	20030709									
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AT 391781 AT 406383	T	T 20080415 AT 2003-763764 2											
ES 2304146	T3	20080915	ES 2003-763764	20030709 20030709									
EP 1995255	A1	20081126	EP 2008-14706	20030709									
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US 20060167227	A1	20060727	ES 2003-763763 US 2005-521140	20051031									
PRAI AT 2002-1053	A	20020712											
PRAI AT 2002-1053 EP 2003-763763 WO 2003-EP7389	A3	20030709											
CLASS	ve	20030709											
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WO 2004007547 ICM ICS		C07K014-47 C12N015-12; G01N033-53; A01K067-027; A61K039-00;											
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AU 2003253044 IPC	I C07K001	C07K0014-47 [ICM, 7]; C07K0014-435 [ICM, 7, C*];											

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EP 1521774
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                       4H045/CA45; 4H045/DA00; 4H045/DA76; 4H045/EA20;
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AB
     Described are novel N- and C-terminally double truncated
     tau mols., (type IA, IB, IIA and IIB tau mols.) as well
     as methods for providing these mols., both from recombinant and biol.
     sources. Moreover, screening methods using these mols. in connection with
     Alzheimer's diagnosis and therapy are provided.
     tau protein fragment isolation sequence Alzheimer diagnosis
     monoclonal antibody
     Transformation, genetic
        (Alzheimer's disease model; N- and C-terminally truncated
        tau proteins isolation and characterization for diagnostic,
        therapeutic and Alzheimer's disease model uses)
     Disease models
        (Alzheimer's disease; N- and C-terminally truncated
        tau proteins isolation and characterization for diagnostic,
        therapeutic and Alzheimer's disease model uses)
     Vaccines
        (Alzheimer's disease; truncated Tau proteins of
        humans)
     Tau factor
     RL: BSU (Biological study, unclassified); DGN (Diagnostic use); PRP
     (Properties); BIOL (Biological study); USES (Uses)
        (N- and C-terminally double truncated; N- and C-terminally
        truncated tau proteins isolation and characterization
        for diagnostic, therapeutic and Alzheimer's disease model uses)
     Alzheimer's disease
     Anti-Alzheimer's agents
     Biomarkers
     Brain
     Diagnosis
     Human
     Neuron
     Oxidative stress, biological
        (N- and C-terminally truncated tau proteins
        isolation and characterization for diagnostic, therapeutic and
       Alzheimer's disease model uses)
     Synthetic gene
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
        (animal; for truncated Tau proteins of humans)
    Microtubule
        (assembly; N- and C-terminally truncated tau
        proteins isolation and characterization for diagnostic, therapeutic and
       Alzheimer's disease model uses)
   Protein sequences
```

(for truncated Tau proteins of humans)

Antibodies and Immunoglobulins

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (monoclonal, DC44, DC82, DC136; N- and C-terminally truncated tau proteins isolation and characterization for diagnostic. therapeutic and Alzheimer's disease model uses)

Gene, animal

RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses)

(synthetic; for truncated Tau proteins of humans)

Antibodies and Immunoglobulins

RL: BSU (Biological study, unclassified); BIOL (Biological study) (to Alzheimer disease derived tau; N- and C-terminally truncated tau proteins isolation and characterization

for diagnostic, therapeutic and Alzheimer's disease model uses) Transferrins

RL: BSU (Biological study, unclassified); DGN (Diagnostic use); PRP (Properties); BIOL (Biological study); USES (Uses) (τ -transferrins; N- and C-terminally truncated

tau proteins isolation and characterization for diagnostic, therapeutic and Alzheimer's disease model uses)

649163-53-1, 239-333-Tau factor (human type IA) 649163-54-2, 237-333-Tau factor (human type IA) 649163-55-3, 239-318-Tau factor (human type IA) 649163-56-4, 239-326-Tau factor (human type IB) 649163-57-5, 239-328-Tau factor (human type IB) 649163-58-6, 239-331-Tau factor (human type IB) 649163-59-7, 239-334-Tau factor (human type IB) 649163-60-0, 239-340-Tau factor (human type IB) 649163-61-1, 239-343-Tau factor (human type IB) 649163-62-2, 208-302-Tau factor (human type IB) 649163-63-3, 69-333-Tau factor (human type IIA) 649163-64-4, 93-333-Tau factor (human type IIA) 649163-65-5, 69-363-Tau factor (human type IIA) 649163-66-6, 93-363-Tau factor (human type IIA) 649163-67-7, 93-302-Tau factor (human type IIA) 649163-68-8, 69-302-Tau factor (human type IIA) 649163-69-9, 93-332-Tau factor (human type IIA) 649163-70-2, 69-332-Tau factor (human type IIA) 649163-71-3, 6-378-Tau factor (human type IIB) 649163-72-4, 6-347-Tau factor (human type IIB) 649163-73-5, 268-333-Tau factor (human) 649163-74-6, 248-333-Tau factor 649163-75-7, 258-333-Tau factor (human) 649163-76-8, (human) 263-333-Tau factor (human) 649163-77-9, (239-247)-(263-333)-Tau factor (human) 649163-78-0, (239-255)-(263-333)-Tau

factor (human) 649163-79-1, (239-262)-(268-333)-Tau factor (human) RL: BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study)

(amino acid sequence; N- and C-terminally truncated tau proteins isolation and characterization for diagnostic. therapeutic and Alzheimer's disease model uses)

RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD RE

- (1) Anon; WO 0118546 A2 CAPLUS
- (2) Anon; WO 0165252 A1 CAPLUS
- (3) Anon: WO 02055720 A2 CAPLUS (4) Anon; WO 02059150 A2 CAPLUS
- (5) Anon; WO 02062851 A1 CAPLUS (6) Anon; WO 9630766 A1 CAPLUS

ENTRY SESSION FULL ESTIMATED COST 24.70 25.36

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1 649163-70-2/RN 1 649163-71-3/RN

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=> s 14 L5 1 L4

=> file uspatfull
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DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

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FILE COVERS 1971 TO PATENT PUBLICATION DATE: 26 Mar 2009 (20090326/PD)
FILE LAST UPDATED: 26 Mar 2009 (20090326/BD)
HIGHEST GRANTED PATENT NUMBER: US7509687
HIGHEST APPLICATION PUBLICATION NUMBER: US20090083889
CA INDEXING IS CURRENT THROUGH 26 Mar 2009 (20090326/UPCA)
ISSUE CLASS FIELDS ((TINCL) CURRENT THROUGH: 26 Mar 2009 (20090326/PD)

REVISED CLASS FIELDS (/NCL) LAST RELOADED: Dec 2008 USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Dec 2008

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L3 4 S L2 AND TRUNCATED

## SELECT RN L3 2

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8 S 649163-64-4/RN OR 649163-65-5/RN OR 649163-66-6/RN OR 6491 T. 4

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TOTAL

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TOTAL.

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L6

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4 S L2 AND TRUNCATED SELECT RN L3 2

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L6

L5 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2004:60551 CAPLUS DOCUMENT NUMBER: 140:124832

TITLE: Truncated tau proteins

INVENTOR(S): Kontsekova, Eva

PATENT ASSIGNEE(S): Axon Neuroscience Forschungs- und Entwicklungs GmbH, Austria

SOURCE: PCT Int. Appl., 97 pp.

CODEN: PIXXD2
DOCUMENT TYPE: Patent

LANGUAGE: English FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

								APPLICATION NO.										
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									EP 2003-763763 WO 2003-EP7389				A3 20030709					
	_										WO 2	2003-1	EP73	89	1	n 2	0030	709

AB Described are novel N- and C-terminally double truncated tau mols., (type IA, IB, IIA and IIB tau mols.) as well as methods for providing these mols., both from recombinant and biol. sources. Moreover, screening methods using these mols. in connection with Alzheimer's diagnosis and

therapy are provided.
REFERENCE COUNT: 6

THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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L2 10 S L1 AND TAU
L3 4 S L2 AND TRUNCATED
SELECT RN L3 2

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FILE 'CAPLUS' ENTERED AT 15:11:11 ON 28 MAR 2009 L5 1 S L4

FILE 'USPATFULL' ENTERED AT 15:11:23 ON 28 MAR 2009 L6 0 S L4

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MOST RECENT UPDATE: 200918 <200918/DW>

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L2 10 S L1 AND TAU
L3 4 S L2 AND TRUNCATED

L3 4 S L2 AND TRUNCATED SELECT RN L3 2

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L4 8 S 649163-64-4/RN OR 649163-65-5/RN OR 649163-66-6/RN OR 6491

FILE 'CAPLUS' ENTERED AT 15:11:11 ON 28 MAR 2009

FILE 'USPATFULL' ENTERED AT 15:11:23 ON 28 MAR 2009 L6  $$0\ \mathrm{S}\ \mathrm{L4}$$ 

FILE 'CAPLUS' ENTERED AT 15:11:57 ON 28 MAR 2009

FILE 'WPIDS' ENTERED AT 15:13:48 ON 28 MAR 2009

L8 0 S L5

FILE 'REGISTRY' ENTERED AT 15:14:33 ON 28 MAR 2009

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L4 ANSWER 1 OF 8 REGISTRY COPYRIGHT 2009 ACS on STN

RN 649163-71-3 REGISTRY

ED Entered STN: 11 Feb 2004

CN 6-378-Tau factor (human type IIB) (9CI) (CA INDEX NAME)

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CN 19: PN: WO2004007547 SEQID: 19 claimed protein
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SR
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LC
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            10 S L1 AND TAU
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FILE 'USPATFULL' ENTERED AT 15:11:23 ON 28 MAR 2009 L6  $$0\ \mathrm{S}\ \mathrm{L4}$$ 

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FILE 'WPIDS' ENTERED AT 15:13:48 ON 28 MAR 2009

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L4